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Application Number		10565308
Filing Date		2006-03-27
First Named Inventor Thorn		as Helleday
Art Unit		1614
Examiner Name	Not Y	et Known
Attorney Docket Number		J660-065 US

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	2	20050227919	A1	2005-10	-13	Ashworth, et a	L			
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	1	2005012524	wo	A1	2005-02-10	The University of Sheffield, et al.		
	2	2004087713	wo	A1	2004-10-14	Pfizer, Inc., et al.		
	3	0116136	wo	A2	2001-03-08	Agouron Pharmaceuticals, Inc., et al.		
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Examiner Initials*	Cite No						Ţ5	
	BRYANT, et al., Specific killing of BRCA2-deficient tumours with inhibitors of poly(ADP-ribose) polymerase, Nature, April 14, 2005, 913-917, Vol. 434.							
	CANAN KOCH, et al., Novel Tricyclic Poly/ADP-rbose) Polymerase-1 Inhibitors with Potent Articanoer Chemopotenisting Activity: Design, Synthesis, and X-ray Coorystal Structure, J. Med. Chem., 2002, 4961-4974, Vol. 45, No. 23.							
	3 EASTON, et al., Carnoer Resks is BRCA2 Mutation Carriers, J. Natl. Canoer Inst., 1999, 1310-1316, Vol. 91, No. 15							
	FARMER, et al., Targeting the DNA repair defect in BRCA mutant cells as a therapeutic strategy, Nature, April 14, 2005, 917-921, Vol. 434							
	5 FRIEDENSON, et al., BRCA1 and BRCA2 Pathways and the Risk of Canoers Other Than Breast or Ovarian, Medicape General Medicine, 2005, 7(2):60.							

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	6	LAXIFABIL of al., The Pathology of Familiat Brosst Cornors: Predictive Value of Immunohistochemical Markers Estrogen Receptor: Progestionene Recognit HERV2 as epid an application With Mutations in BRCA1 and BRCA2, Journal of Clinical Chrology, May 1, 2002, 2310-2318, Vol. 29, No.						
	7	QUINN, et al., BRCA1 Functions as a Different Modulator of Chemotherapy-induced Apoptosis, Cancer Research, October 1, 2003, 6221-6228, Vol. 63.						
	8	TARON, et al., BRCA1 mRNA expression levels as an indicator of chemoresistance in lung cancer, Human Molecular Genetics, 2004, 2443-2449, Vol. 13, No. 20.						
	9	TUTT, et al., The relationship between the roles of BRCA genes in DNA repair and cancer predisposition, Trends in Molecular Medicine, December 2002, 571-576, Vol. 8, No. 12.						
	10	VENKITARAMAN, A. R., Cancer Susceptibility and the Functions of BRCA1 and BRCA2, Cell, January 25, 2002, 171-182, Vol. 108.						
	11	PCT international Search Report for PCT/GB2004/003183 (WO 2005/012305).						
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That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patient office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. Sea 37 CFF 1.37(e)(1).

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A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Chih-Sheng Lin/	Date (YYYY-MM-DD)	2007-01-10
Name/Brint	Ohib Ohana Lin	Registration Number	66402

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